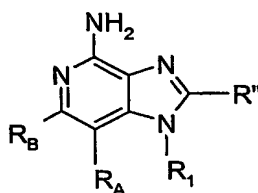


WHAT IS CLAIMED IS:

1. A compound of the following Formula I:



I

wherein:

- 10 R_1 has the formula alkylene-L- R_{1-1} , alkenylene-L- R_{1-1} , or alkynylene-L- R_{1-1} , wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group; and

- 15 R_{1-1} is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

R'' is hydrogen or a non-interfering substituent;

R_A and R_B are each independently selected from the group consisting of:

- 20 hydrogen,
halogen,
alkyl,
alkenyl,
alkoxy,
alkylthio, and
25 $-N(R_3)_2$;

- or when taken together, R_A and R_B form a fused aryl ring or heteroaryl ring containing one heteroatom or a fused 5- to 7-membered saturated ring, optionally containing one heteroatom, wherein the heteroatom is selected from the group consisting of N and S, and wherein the aryl, heteroaryl, or 5- to 7-membered saturated ring is unsubstituted or substituted by one or more non-interfering substituents; and
- 30

each R_3 is independently selected from the group consisting of hydrogen and alkyl;

with the proviso that when L is $-NH-S(O)_2-$ and R_A and R_B join to form an unsubstituted benzene ring, R_{1-1} is a linear or branched aliphatic group having greater than 16 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds; and with the further proviso that when L is $-NH-C(O)-$ and R_A and R_B join to form an unsubstituted pyridine ring, R_{1-1} is a linear or branched aliphatic group having greater than 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds; or a pharmaceutically acceptable salt thereof.

2. The compound or salt of claim 1 wherein when taken together, R_A and R_B form a fused 5- to 7-membered saturated ring, optionally containing one heteroatom selected from the group consisting of N and S, and unsubstituted or substituted by one or more substituents selected from the group consisting of:

halogen,
hydroxy,
alkyl,
alkenyl,
haloalkyl,
alkoxy,
alkylthio, and
 $-N(R_3)_2$.

3. The compound or salt of claim 1 wherein R_A and R_B are each independently selected from the group consisting of:

hydrogen,
halogen,
alkyl,
alkenyl,

alkoxy,
alkylthio and
-N(R₃)₂.

5 4. The compound or salt of claim 1 wherein R_A and R_B form a fused aryl or heteroaryl ring.

5. The compound or salt of claim 1 wherein R_A and R_B form a fused 5- to 7-membered saturated ring.

10

6. The compound or salt of claim 1 wherein when taken together, R_A and R_B form a fused aryl ring or heteroaryl ring containing one heteroatom selected from the group consisting of N and S wherein the aryl or heteroaryl ring is unsubstituted or substituted by one or more R groups;

15 or when taken together, R_A and R_B form a fused 5- to 7-membered saturated ring, optionally containing one heteroatom selected from the group consisting of N and S, and unsubstituted or substituted by one or more R groups;

each R is independently selected from the group consisting of

20 halogen,
hydroxy,
alkyl,
alkenyl,
haloalkyl,
alkoxy,
25 alkylthio, and
-N(R₃)₂.

7. The compound or salt of claim 6 wherein R_A and R_B form a fused benzene ring which is unsubstituted.

30

8. The compound or salt of claim 6 wherein R_A and R_B form a fused pyridine ring which is unsubstituted.

9. The compound or salt of any one of claims 1 through 8 wherein R" is selected from the group consisting of:

- hydrogen;
- alkyl;
- 5 alkenyl;
- aryl;
- heteroaryl;
- heterocyclyl;
- alkylene-Y-alkyl;
- 10 alkylene-Y- alkenyl;
- alkylene-Y-aryl; and
- alkyl or alkenyl substituted by one or more substituents selected

from the group consisting of:

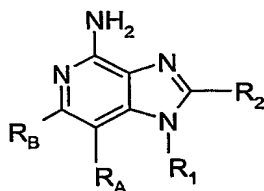
- OH;
- 15 halogen;
- N(R₄)₂;
- C(O)-C₁₋₁₀alkyl;
- C(O)-O-C₁₋₁₀alkyl;
- N₃;
- 20 aryl;
- heteroaryl;
- heterocyclyl;
- C(O)-aryl; and
- C(O)-heteroaryl;

25 wherein: Y is -O- or -S(O)₀₋₂-; and each R₄ is independently selected from the group consisting of hydrogen, C₁₋₁₀alkyl, and C₂₋₁₀alkenyl.

10. The compound or salt of any one of claims 1 through 9 wherein L is a bond or a functional linking group selected from the group consisting of

30 -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-, -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-,
 -NH-C(S)-NR₃-, -NH-C(O)-O-, -O-, -S-, and -S(O)₂-.

11. The compound or salt of claim 10 wherein L is a bond or a functional linking group selected from the group consisting of -NH-C(O)-, -NH-S(O)₂-, and -NH-C(O)-N(R₃)-.
 5 12. The compound or salt of any one of claims 1 through 11 wherein R₁₋₁ is a linear or branched aliphatic group having 11-20 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds.
 10 13. The compound or salt of claim 12 wherein R₁₋₁ is a linear or branched aliphatic group having 12-20 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds.
 14. The compound or salt of claim 13 wherein R₁₋₁ is a straight chain C₁₂-C₂₀alkyl.
 15 15. A compound of the following Formula II:



II

wherein:

R₁ has the formula alkylene-L-R₁₋₁, alkenylene-L-R₁₋₁, or alkynylene-L-R₁₋₁, wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group selected from the group consisting of -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-, -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-, -NH-C(S)-NR₃-, -NH-C(O)-O-, -O-, -S-, and -S(O)₂-; and

R_{1-1} is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

R_2 is selected from the group consisting of:

- 5 hydrogen;
- alkyl;
- alkenyl;
- aryl;
- heteroaryl;
- 10 heterocyclyl;
- alkylene-Y-alkyl;
- alkylene-Y-alkenyl;
- alkylene-Y-aryl; and
- alkyl or alkenyl substituted by one or more substituents selected

15 from the group consisting of:

- OH;
- halogen;
- N(R₄)₂;
- C(O)-C₁₋₁₀alkyl;
- 20 -C(O)-O-C₁₋₁₀alkyl;
- N₃;
- aryl;
- heteroaryl;
- heterocyclyl;
- 25 -C(O)-aryl; and
- C(O)-heteroaryl;

wherein: Y is -O- or -S(O)₀₋₂-; and each R₄ is independently selected from the group consisting of hydrogen, C₁₋₁₀alkyl, and C₂₋₁₀alkenyl;

30 R_A and R_B are each independently selected from the group consisting of:

- hydrogen,
- halogen,
- alkyl,

alkenyl,
alkoxy,
alkylthio, and
-N(R₃)₂;

5 or when taken together, R_A and R_B form a fused aryl ring or heteroaryl ring containing one heteroatom wherein the aryl or heteroaryl ring is unsubstituted or substituted by one or more R groups; or when taken together, R_A and R_B form a fused 5- to 7-membered saturated ring, optionally containing one heteroatom selected from the group consisting of N and S, and unsubstituted or
10 substituted by one or more R groups; wherein R is selected from the group consisting of

halogen,
hydroxy,
alkyl,
15 alkenyl,
haloalkyl,
alkoxy,
alkylthio, and
-N(R₃)₂.

20 and

R₃ is selected from the group consisting of hydrogen and alkyl; with the proviso that when L is -NH-S(O₂)- and R_A and R_B join to form an unsubstituted benzene ring, R₁₋₁ is a linear or branched aliphatic group having at least 16 carbon atoms, optionally including one or more unsaturated carbon-
25 carbon bonds; and with the further proviso that when L is -NH-C(O)- and R_A and R_B join to form an unsubstituted pyridine ring, R₁₋₁ is a linear or branched aliphatic group having greater than 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;
or a pharmaceutically acceptable salt thereof.

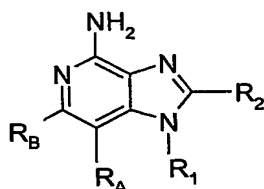
30

16. The compound or salt of claim 15 wherein R₁ has the formula alkylene-L-R₁₋₁ and the alkylene is optionally interrupted with one -O- group.

17. The compound or salt of claim 16 wherein R_1 has the formula C_{1-5} alkylene-L- R_{1-1} and the C_{1-5} alkylene is optionally interrupted with one -O- group.

5 18. The compound or salt of claim 15 wherein R_2 is selected from the group consisting of hydrogen, alkyl, and alkylene-O-alkyl.

19. A compound of the following Formula II:



10

II

wherein:

15 R_1 has the formula alkylene-L- R_{1-1} , alkenylene-L- R_{1-1} , or alkynylene-L- R_{1-1} , wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

20 L is a bond or a functional linking group selected from the group consisting of -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-, -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-, -NH-C(S)-NR₃-, -NH-C(O)-O-, -O-, -S-, and -S(O)₂-; and

R_{1-1} is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

25 R_2 is selected from the group consisting of:

hydrogen;

alkyl;

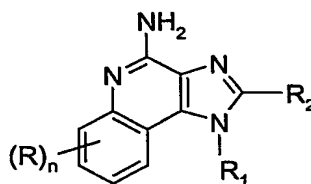
alkenyl;

aryl;

30 heteroaryl;

- heterocyclyl;
alkylene-Y-alkyl;
alkylene-Y-alkenyl;
alkylene-Y-aryl; and
5 alkyl or alkenyl substituted by one or more substituents selected
from the group consisting of:
-OH;
halogen;
-N(R₄)₂;
10 -C(O)-C₁₋₁₀alkyl;
-C(O)-O-C₁₋₁₀alkyl;
-N₃;
aryl;
heteroaryl;
15 heterocyclyl;
-C(O)-aryl; and
-C(O)-heteroaryl;
wherein: Y is -O- or -S(O)₀₋₂-; and each R₄ is
independently selected from the group consisting of hydrogen,
20 C₁₋₁₀alkyl, and C₂₋₁₀alkenyl;
R_A and R_B are each independently selected from the group consisting of:
hydrogen,
halogen,
alkyl,
25 alkenyl,
alkoxy,
alkylthio, and
-N(R₃)₂; and
R₃ is selected from the group consisting of hydrogen and alkyl;
30 or a pharmaceutically acceptable salt thereof.

20. A compound of the following Formula III:



III

wherein:

R_1 has the formula alkylene-L- R_{1-1} , alkenylene-L- R_{1-1} , or
 5 alkynylene-L- R_{1-1} , wherein:

the alkylene, alkenylene, and alkynylene groups are
 optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group selected from the
 group consisting of -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-,
 10 -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-, -NH-C(S)-NR₃-,
 -NH-C(O)-O-, -O-, -S-, and -S(O)₂-; and

R_{1-1} is a linear or branched aliphatic group having at least
 11 carbon atoms, optionally including one or more unsaturated
 carbon-carbon bonds;

15 R is selected from the group consisting of

halogen,
 hydroxy,
 alkyl,
 alkenyl,
 20 haloalkyl,
 alkoxy,
 alkylthio, and
 -N(R₃)₂;

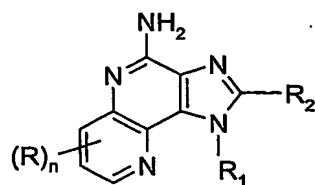
n is 0 to 4;

25 R_2 is selected from the group consisting of:

hydrogen;
 alkyl;
 alkenyl;
 aryl;
 30 heteroaryl;

- heterocyclyl;
alkylene-Y-alkyl;
alkylene-Y-alkenyl;
alkylene-Y-aryl; and
5 alkyl or alkenyl substituted by one or more substituents selected
from the group consisting of:
-OH;
halogen;
-N(R₄)₂;
10 -C(O)-C₁₋₁₀alkyl;
-C(O)-O-C₁₋₁₀alkyl;
-N₃;
aryl;
heteroaryl;
15 heterocyclyl;
-C(O)-aryl; and
-C(O)-heteroaryl;
Y is -O- or -S(O)₀₋₂;
each R₄ is independently selected from the group consisting of hydrogen,
20 C₁₋₁₀alkyl, and C₂₋₁₀alkenyl; and
R₃ is selected from the group consisting of hydrogen and alkyl;
with the proviso that when L is -NH-S(O₂)-, and n is 0, R₁₋₁ is a linear or
branched aliphatic group having at least 16 carbon atoms, optionally including
one or more unsaturated carbon-carbon bonds;
25 or a pharmaceutically acceptable salt thereof.

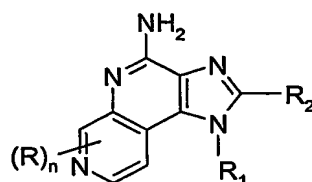
21. The compound or salt of claim 20 wherein n is 0.
22. A compound selected from the group consisting of the following
30 Formulas IV, V, VI, and VII:



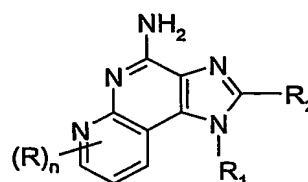
IV



V



VI



VII

5

wherein:

R_1 has the formula alkylene-L- R_{1-1} , alkenylene-L- R_{1-1} , or alkynylene-L- R_{1-1} , wherein:

the alkylene, alkenylene, and alkynylene groups are optionally interrupted with one or more -O- groups;

L is a bond or a functional linking group selected from the group consisting of -NH-S(O)₂-, -NH-C(O)-, -NH-C(S)-, -NH-S(O)₂-NR₃-, -NH-C(O)-NR₃-, -NH-C(S)-NR₃-, -NH-C(O)-O-, -O-, -S-, and -S(O)₂-; and

R_{1-1} is a linear or branched aliphatic group having at least 11 carbon atoms, optionally including one or more unsaturated carbon-carbon bonds;

R is selected from the group consisting of

halogen,
hydroxy,
alkyl,
alkenyl,
haloalkyl,
alkoxy,
alkylthio, and
-N(R₃)₂;

n is 0 or 1;

R₂ is selected from the group consisting of:

- hydrogen;
alkyl;
alkenyl;
5 aryl;
heteroaryl;
heterocyclyl;
alkylene-Y-alkyl;
alkylene-Y-alkenyl;
10 alkylene-Y-aryl; and
alkyl or alkenyl substituted by one or more substituents selected
from the group consisting of:

- OH;
halogen;
15 -N(R₄)₂;
-C(O)-C₁₋₁₀alkyl;
-C(O)-O-C₁₋₁₀alkyl;
-N₃;
aryl;
20 heteroaryl;
heterocyclyl;
-C(O)-aryl; and
-C(O)-heteroaryl;

Y is -O- or -S(O)₀₋₂;

- 25 each R₄ is independently selected from the group consisting of hydrogen,
C₁₋₁₀alkyl, and C₂₋₁₀alkenyl; and

- R₃ is selected from the group consisting of hydrogen and alkyl;
with the proviso that when L is -NH-C(O)-, and n is 0, R₁₋₁ is a linear or
branched aliphatic group having at least 12 carbon atoms, optionally including
30 one or more unsaturated carbon-carbon bonds;
or a pharmaceutically acceptable salt thereof.

23. The compound or salt of claim 22 wherein n is 0.

24. A pharmaceutical composition comprising a therapeutically effective amount of a compound or salt of any one of claims 1 through 23 in combination with a pharmaceutically acceptable carrier.
- 5
25. A method of inducing cytokine biosynthesis in an animal comprising administering an effective amount of a compound or salt of any one of claims 1 through 23 to the animal.
- 10
26. A method of treating a viral disease in an animal comprising administering a therapeutically effective amount of a compound or salt of any one of claims 1 through 23 to the animal.
- 15
27. A method of treating a neoplastic disease in an animal comprising administering a therapeutically effective amount of a compound or salt of any one of claims 1 through 23 to the animal.
- 20
28. A method of vaccinating an animal comprising administering an effective amount of a compound or salt of any one of claims 1 through 23 to the animal as a vaccine adjuvant.
- 25
29. A method of vaccinating an animal comprising administering an effective amount of *N*-(2-{2-[4-amino-2-(2-methoxyethyl)-1*H*-imidazo[4,5-*c*]quinolin-1-yl]ethoxy}ethyl)hexadecanamide to the animal as a vaccine adjuvant.
- 30
30. A method of vaccinating an animal comprising administering an effective amount of *N*-(2-{2-[4-amino-2-(2-methoxyethyl)-1*H*-imidazo[4,5-*c*]quinolin-1-yl]ethoxy}ethyl)octadecanamide to the animal as a vaccine adjuvant.
- 30
31. A method of vaccinating an animal comprising administering an effective amount of *N*-(2-{2-[4-amino-2-(2-methoxyethyl)-1*H*-imidazo[4,5-*c*]quinolin-1-yl]ethoxy}ethyl)dodecanamide to the animal as a vaccine adjuvant.

32. A method of vaccinating an animal comprising administering an effective amount of *N*-(2-{2-[4-amino-2-(2-methoxyethyl)-1*H*-imidazo[4,5-*c*]quinolin-1-yl]ethoxy}ethyl)tetradecanamide to the animal as a vaccine adjuvant.